

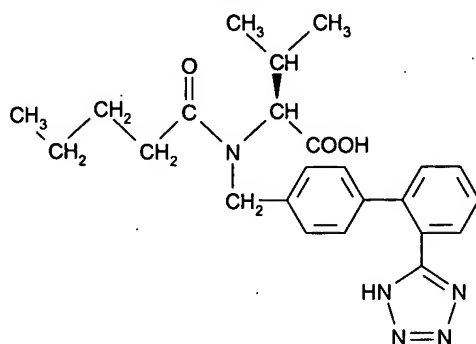
This listing of the claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

Please delete claims 8, 9, 11 and 12.

1-14. (canceled)

15. (withdrawn) A method of treating obstructive airways diseases, wherein the diseases are selected from the group consisting of chronic obstructive pulmonary disease, adult respiratory distress syndrome, sepsis syndrome, pneumonia, aspiration of gastric content, chest trauma, shock, burns, fat embolia, cardiopulmonary bypass, O₂ toxicity, haemorrhagic pancreatitis, interstitial and bronchoalveolar inflammation comprising administering a therapeutically effective amount of valsartan of formula



to a patient in need thereof.

16. (withdrawn) The method of claim 15 wherein chronic obstructive pulmonary disease includes bronchitis, chronic bronchitis, emphysema, asthma, cystic fibrosis, interstitial lung disease, pulmonary vascular disease and increased resistance to airflow during forced expiration.

17. (withdrawn) A solid oral dosage form according to claim 8 comprising 20 to 65% of valsartan.

18. (withdrawn) A solid oral dosage form according to claim 9 comprising 20 to 65% of valsartan.

19 (withdrawn) A solid oral dosage form according to claim 10 comprising 20 to 65% of valsartan.

20. (withdrawn) A solid oral dosage form according to claim 8 comprising 20 to 360 mg of valsartan.

21. (withdrawn) A solid oral dosage form according to claim 9 comprising 20 to 360 mg of valsartan.
22. (withdrawn) A solid oral dosage form according to claim 10 comprising 20 to 360 mg of valsartan.
23. (withdrawn) A solid oral dosage form according to claim 11 comprising 20 to 360 mg of valsartan.
24. (withdrawn) A solid oral dosage form according to claim 8 comprising less than 13% of crospovidone.
25. (new) A compressed tablet comprising valsartan, microcrystalline cellulose and crospovidone, wherein the weight ratio of valsartan to microcrystalline cellulose is from 2:1 to 0.3 : 1.
26. (new) A tablet according to claim 25 wherein the weight ratio of valsartan to microcrystalline cellulose is from 2 : 1 to 1 : 1.
27. (new) A tablet according to claim 25 comprising less than 13 % by weight of crospovidone.
28. (new) A tablet according to any one of claims 25, wherein the weight ratio of microcrystalline cellulose to crospovidone is from 7 : 1 to 2 : 1.
29. (new) A tablet according to any one of claim 25, wherein the weight ratio of microcrystalline cellulose to crospovidone is from 4 : 1 to 2 : 1.
30. (new) A tablet according to claim 25, wherein the tablet comprises
20 to 65 % of valsartan,
31 to 65 % of microcrystalline cellulose and
2 to 13 % of crospovidone.
31. (new) A tablet according to claim 25, wherein the tablet comprises more than 250 mg and up to 360 mg valsartan as an active agent.
32. (new) A tablet according to claim 25, wherein the tablet comprises 320 mg valsartan as an active agent.

33. (new) A tablet according to claim 25, comprising
40 mg of valsartan
27 mg of microcrystalline cellulose and
7.5 mg of crospovidone.

34. (new) A tablet according to claim 25, comprising
80 mg of valsartan
54 mg of microcrystalline cellulose and
15 mg of crospovidone.

35. (new) A tablet according to claim 25 comprising
160 mg of valsartan
108 mg of microcrystalline cellulose and
30 mg of crospovidone.

36. (new) A tablet according to claim 25, comprising
320 mg of valsartan
216 mg of microcrystalline cellulose and
60 mg of crospovidone.

37. (new) A tablet according to claim 25 comprising valsartan in free form.

38. (new) A tablet according to claim 25 having a dissolution rate of above 90 % over 30 minutes.